- (i) A compound that contains at least one ingredient covered under Part B as prescribed and dispensed or administered is considered a Part B compound, regardless of whether other ingredients in the compound are covered under Part B as prescribed and dispensed or administered.
- (ii) Only compounds that contain at least one ingredient that independently meets the definition of a Part D drug, and that do not meet the criteria under paragraph (d)(1)(i) of this section, may be covered under Part D. For purposes of this paragraph (d) these compounds are referred to as Part D compounds.
- (iii) For a Part D compound to be considered on-formulary, all ingredients that independently meet the definition of a Part D drug must be considered on-formulary (even if the particular Part D drug would be considered off-formulary if it were provided separately—that is, not as part of the Part D compound).
- (iv) For a Part D compound that is considered off-formulary—
- (A) Transition rules apply such that all ingredients in the Part D compound that independently meet the definition of a Part D drug must become payable in the event of a transition fill under §423.120(b)(3); and
- (B) All ingredients that independently meet the definition of a Part D drug must be covered if an exception under § 423.578(b) is approved for coverage of the compound.
- (2) Establish consistent rules for beneficiary payment liabilities for both ingredients of the Part D compound that independently meet the definition of a Part D drug and non-Part D ingredients.
- (i) For low income subsidy beneficiaries the copayment amount is based on whether the most expensive ingredient that independently meets the definition of a Part D drug in the Part D compound is a generic or brand name drug (as described under § 423,782).
- (ii) For any non-Part D ingredient of the Part D compound (including drugs described under $\S423.104(f)(1)(ii)(A)$), the Part D sponsor's contract with the pharmacy must prohibit balance bill-

ing the beneficiary for the cost of any such ingredients.

[70 FR 4525, Jan. 28, 2005, as amended at 73 FR 20506, Apr. 15, 2008; 74 FR 2888, Jan. 16, 2009; 75 FR 19816, Apr. 15, 2010; 75 FR 32860, June 10, 2010; 76 FR 21572, Apr. 15, 2011; 77 FR 22169, Apr. 12, 2012]

§ 423.124 Special rules for out-of-network access to covered Part D drugs at out-of-network pharmacies.

- (a) Out-of-network access to covered part D drugs—(1) Out-of-network pharmacy access. A Part D sponsor must ensure that Part D enrollees have adequate access to covered Part D drugs dispensed at out-of-network pharmacies when the enrollees—
- (i) Cannot reasonably be expected to obtain such drugs at a network pharmacy; and
- (ii) Do not access covered Part D drugs at an out-of-network pharmacy on a routine basis.
- (2) Physician's office access. A Part D sponsor must ensure that Part D enrollees have adequate access to vaccines and other covered Part D drugs appropriately dispensed and administered by a physician in a physician's office.
- (b) Financial responsibility for out-ofnetwork access to covered Part D drugs. A
 Part D sponsor that provides its Part D
 enrollees with coverage other than defined standard coverage may require
 its Part D enrollees accessing covered
 Part D drugs as provided in paragraph
 (a) of this section to assume financial
 responsibility for any differential between the out-of-network pharmacy's
 (or provider's) usual and customary
 price and the Part D sponsor's plan allowance, consistent with the requirements of §§ 423.104(d)(2)(i)(B) and
 423.104(e).
- (c) Limits on out-of-network access to covered Part D. A Part D sponsor must establish reasonable rules to appropriately limit out-of-network access to covered Part D drugs.

§ 423.128 Dissemination of Part D plan information.

(a) Detailed description. A Part D sponsor must disclose the information specified in paragraph (b) of this section in the manner specified by CMS—